

Regimen Monograph

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A - Regimen Name

CAPEMTMC(RT) Regimen

Capecitabine-mitomycin

Disease Site Genitourinary
Penile

Intent Adjuvant
Curative

Regimen Category **Evidence-informed :**
Regimen is considered appropriate as part of the standard care of patients; meaningfully improves outcomes (survival, quality of life), tolerability or costs compared to alternatives (recommended by the Disease Site Team and national consensus body e.g. pan-Canadian Oncology Drug Review, pCODR). Recommendation is based on an appropriately conducted phase III clinical trial relevant to the Canadian context OR (where phase III trials are not feasible) an appropriately sized phase II trial. Regimens where one or more drugs are not approved by Health Canada for any indication will be identified under Rationale and Use.

This **Regimen Abstract** is an **abbreviated** version of a Regimen Monograph and contains only top level information on usage, dosing, schedule, cycle length and special notes (if available). Information in regimen abstracts is accurate to the extent of the ST-QBP regimen master listings, and has not undergone the full review process of a regimen monograph. Full regimen monographs will be published for each ST-QBP regimen as they are developed.

Rationale and Uses For treatment of locally advanced penile cancer

Supplementary Public Funding [capecitabine](#)
ODB - General Benefit (capecitabine)

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| | | | |
|---------------------|------------------------|----|------------------------|
| <u>capecitabine</u> | 825 mg /m ² | PO | BID* on radiation days |
|---------------------|------------------------|----|------------------------|

*total daily dose 1650 mg/m²

| | | | |
|------------------|-----------------------|-------------------|---------------|
| <u>mitomycin</u> | 10 mg /m ² | IV (max 15-20 mg) | Days 1 and 29 |
|------------------|-----------------------|-------------------|---------------|

Alternative mitomycin schedule:

| | | | |
|------------------|-----------------------------|-------------------|------------|
| <u>mitomycin</u> | 10 or 12 mg /m ² | IV (max 15-20 mg) | Day 1 ONLY |
|------------------|-----------------------------|-------------------|------------|

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Single course, concurrent with radiotherapy

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Antiemetic Regimen: Low – No routine prophylaxis; PRN recommended

- Also refer to [CCO Antiemetic Recommendations](#).

Screen for hepatitis B virus in all cancer patients starting systemic treatment. Refer to the [hepatitis B virus screening and management](#) guideline.

Other Supportive Care:

- Topical emollients (e.g. hand creams, udder balm) may ameliorate the manifestations of hand-foot syndrome in patients receiving capecitabine.
- Supportive care should be provided, including loperamide for diarrhea.

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Outpatient prescription for home administration (capecitabine)

| | |
|--|----------------|
| Approximate Patient Visit | 0.5 hour |
| Pharmacy Workload (average time per visit) | 16.99 minutes |
| Nursing Workload (average time per visit) | 41.667 minutes |

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Ajani JA, Winter KA, Gunderson LL, et al. Fluorouracil, mitomycin, and radiotherapy vs fluorouracil, cisplatin, and radiotherapy for carcinoma of the anal canal: a randomized trial. *JAMA* 2008;299:1914-21.

Glynne-Jones R, Meadows H, Wan S, et al. EXTRA--a multicenter phase II study of chemoradiation using a 5 day per week oral regimen of capecitabine and intravenous mitomycin C in anal cancer. *Int J Radiat Oncol Biol Phys*. 2008 Sep 1;72(1):119-26.

Meulendijks D, Dewit L, Tomasoa NB, et al. Chemoradiotherapy with capecitabine for locally advanced anal carcinoma: an alternative treatment option. *Br J Cancer*. 2014 Oct 28;111(9):1726-33.

NCCN clinical practice guidelines in oncology: Penile cancer (v1.2026). November 12, 2025.

Ottenhof SR, de Vries HM, Doodeman B, et al. A prospective study of chemoradiotherapy as primary treatment in patients with locoregionally advanced penile carcinoma. *Int J Radiat Oncol Biol Phys*. 2023 Sep 1;117(1):139-47.

Richter S, Ruether JD, Wood L, et al. Management of carcinoma of the penis: Consensus statement from the Canadian Association of Genitourinary Medical Oncologists (CAGMO). *Can Urol Assoc J*. 2013 Nov-Dec;7(11-12):E797-811.

Thind G, Johal B, Follwell M, Kennecke HF. Chemoradiation with capecitabine and mitomycin-C for stage I-III anal squamous cell carcinoma. *Radiat Oncol*. 2014 May 29;9:124.

PEBC Advice Documents or Guidelines

- [An Endorsement of the 2023 European Association of Urology \(EAU\) -American Society of Clinical Oncology \(ASCO\) Guidelines on Penile Cancer](#)

January 2026 new ST-QBP regimen

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L - Other Notes

DPD Deficiency Testing and Guidance:

Patients should be tested for DPD deficiency before starting treatment with capecitabine. Refer to the [DPD Deficiency Guidance for Clinicians](#) for more information.

In patients with unrecognized DPD deficiency, acute, life-threatening toxicity may occur; if acute grade 2-4 toxicity develops, treatment should be stopped immediately and permanent discontinuation considered based on clinical assessment of the toxicities.

Antidote for Capecitabine Overdose:

Uridine triacetate is a prodrug of uridine and is a specific antidote for treating capecitabine overdose or severe early onset toxicities. If available, consider administering as soon as possible (i.e. within 96 hours) for suspected overdose. If not available, treatment is symptomatic and supportive.

For usage approval and supply, contact Health Canada's [Special Access Program](#) (SAP) (Phone: 613-941-2108. On-call service is available for emergencies).

The recommended dosing and administration for **uridine triacetate** in patients ≥ 18 years is:

- 10 grams (1 packet of coated granules) orally every 6 hours for 20 doses in total, without regards to meals.
- Granules should not be chewed. They should be mixed with 3 to 4 ounces of soft foods such as applesauce, pudding or yogurt.
- The dose should be ingested within 30 minutes of preparation, followed by at least 4 ounces of water.
- Refer to the prescribing information on dose preparation for NG-tube or G-tube use.

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M - Disclaimer

Regimen Abstracts

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Regimen Monographs

Refer to the [New Drug Funding Program](#) or [Ontario Public Drug Programs](#) websites for the most up-to-date public funding information.

The information set out in the drug monographs, regimen monographs, appendices and symptom management information (for health professionals) contained in the *Drug Formulary* (the "Formulary") is intended for healthcare providers and is to be used for informational purposes only. The information is not intended to cover all possible uses, directions, precautions, drug interactions or adverse effects of a particular drug, nor should it be construed to indicate that use of a particular drug is safe, appropriate or effective for a given condition. The information in the Formulary is not intended to constitute or be a substitute for medical advice and should not be relied upon in any such regard. All uses of the Formulary are subject to clinical judgment and actual prescribing patterns may not follow the information provided in the Formulary.

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Some Formulary documents, such as the medication information sheets, regimen information sheets and symptom management information (for patients), are intended for patients. Patients should always consult with their healthcare provider if they have questions regarding any information set out in the Formulary documents.

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